In the Claims

- 1-10. (Canceled)
- 11. (Currently amended) An implantable device comprising a coating, which comprises a polymeric composition comprising a polysulfone (A) and an elastomeric polymer (B), wherein the polysulfone and the elastomeric polymer form a conjugate is selected from the group consisting of polyisobutylene, polyperfluoroalkylene, polyhexafluoropentene, polysiloxane, natural rubber, nylon, polymers and copolymers of acrylates or methacrylates and a combination thereof.
- 12. (Previously presented) The implantable device of claim 11, wherein the device is a stent.
- 13. (Previously presented) The implantable device of claim 12, wherein the coating further comprises a bioactive agent.
- 14. (Previously presented) The implantable device of claim 13 wherein the bioactive agent is selected from the group consisting of tacrolimus, dexamethasone, rapamycin, Everolimus, 40-O-(3-hydroxy)propyl-rapamycin, 40-O-[2-(2-hydroxy)ethoxy]ethyl-rapamycin, and 40-O-tetrazole-rapamycin, paclitaxel, taxoids, estradiol, steroidal anti-inflammatory agents, antibiotics, nitric oxide donors, super oxide dismutases, super oxide dismutase mimics, 4-amino-2,2,6,6-tetramethylpiperidine-1-oxyl (4-amino-TEMPO), and a combination thereof.
 - 15-30. (Canceled)
 - 31. (Canceled)
- 32. (Currently amended) The implantable device of claim 11, wherein the polysulfone and the elastomeric polymer form a conjugate is selected from the group consisting of polyisobutylene, polyperfluoroalkylene, polyhexafluoropentene, polysiloxane, natural rubber, nylon, poly(butyl methacrylate), poly(lauryl methacrylate), polyalkylene oxide, polyalkylene

oxide acrylate, and a combination thereof.

- 33. (Previously presented) The implantable device of claim 11, wherein the polysulfone and the elastomeric polymer form a polymer blend.
- 34. (Currently amended) The implantable device of claim 3211, wherein the conjugate comprises a copolymer that comprises at least one block of a polysulfone polymer (A) and at least one block of an elastomeric polymer (B) in a general formula selected from AB, ABA or BAB.
- 35. (Currently amended) The implantable device of claim 34, wherein the block copolymer is selected from the group consisting of

wherein R₁ is selected from the group consisting of C1 to C10 alkyl, C2, C4 and C6 hydroxyalkyl, C1 to C6 fluoroalkyl, phenyl, substituted phenyl, polyethylene glycol, polyalkylene oxide, ethylene oxide and propylene oxide;

wherein R₂, R₄, R₅, R₇ and R₈ are independently selected from the group consisting of hydrogen, C1 to C6 alkyl, C2, C4 and C6 hydroxyalkyl, C1 to C6 fluoroalkyl, phenyl, substituted phenyl, carboxyl, amido, ester groups bearing a polyethylene glycol, and polyalkylene oxide;

wherein R₃ is selected from the group consisting of hydrogen, alkyl, cycloalkyl, phenyl, carboxyl, halo, amino, hydroxyl, amido, sulfido, and polyalkylene oxide;

wherein R₆ is a perfluoroalkyl group;

wherein
$$R_4$$
, R_5 , and R_6 are selected such that the

$$\begin{bmatrix}
F_2 & | \\
C & C
\end{bmatrix}_{m \text{ are elastomeric; and}}$$

wherein n and m are independently positive integers.

36. (Currently amended) The implantable device of claim 35, wherein R_1 is butyl, isobutyl or isopropyl;

wherein R₂ is hydrogen or methyl;

wherein R₃ is hydrogen, halo, or methyl;

wherein R_4 and R_5 are independently hydrogen, methyl, ethyl, isopropyl, butyl, isobutyl, or phenyl;

wherein R_6 is F, CF_3 , CF_2CF_3 , CF_2CF_3 , perfluoroisopropyl, perfluorobutyl or perfluoroisobutyl; and

wherein R_7 and R_8 are independently methyl, ethyl, propyl, isopropyl, butyl, or isobutyl group.

37. (Currently amended) The implantable device of claim 35, wherein R_1 is butyl;

wherein R₂ is methyl;

wherein R₃ is hydrogen;

wherein R_4 and or R_5 are methyl groups;

wherein R₆ is CF₃; and

wherein R₇ and R₈ are methyl groups.

- 38. (Canceled)
- 39. (Canceled)